

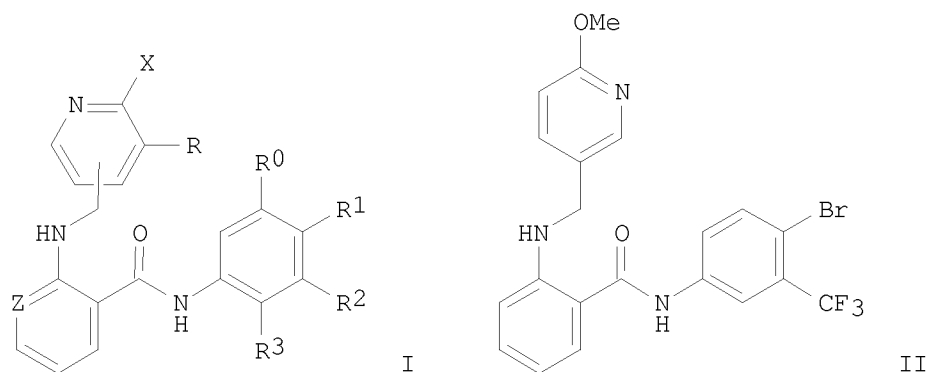
STN Search

L11 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:515506 HCAPLUS
DOCUMENT NUMBER: 141:71453
TITLE: Preparation of anthranilic acid amide derivatives as
neoplastic inhibitors
INVENTOR(S): Bold, Guido; Furet, Pascal; Manley, Paul William
PATENT ASSIGNEE(S): Novartis Ag, Switz.; Novartis Pharma GmbH
SOURCE: PCT Int. Appl., 81 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2004052884	A1	20040624	WO 2003-EP14086	20031211
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LT, LU, LV, MA, MD, MK, MN, MX, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SE, SG, SK, SY, TJ, TM, TN, TR, TT, UA, US, UZ, VC, VN, YU, ZA, ZW				
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CA 2506164	A1	20040624	CA 2003-2506164	20031211
AU 2003294834	A1	20040630	AU 2003-294834	20031211
EP 1572686	A1	20050914	EP 2003-785795	20031211
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BR 2003017292	A	20051108	BR 2003-17292	20031211
CN 1720244	A	20060111	CN 2003-80104845	20031211
JP 2006511518	T	20060406	JP 2004-558075	20031211
US 20060128684	A1	20060615	US 2005-538199	20050609
PRIORITY APPLN. INFO.:			GB 2002-29022	A 20021212
			WO 2003-EP14086	W 20031211
OTHER SOURCE(S):		MARPAT 141:71453		
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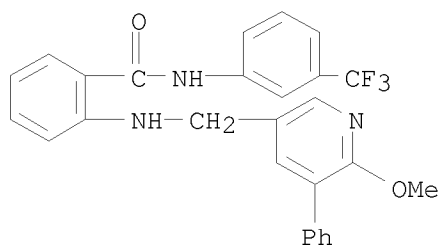


AB The title compds. I [wherein R and R₀ = independently H, halo, (un)substituted alkyl, alkenyl, alkynyl, aryl, heteroaryl, etc.; R₁ = H, halo, (un)substituted alkyl, alkenyl, alkynyl, alkoxy, OCF₃, OCH₂CF₃, OCH₂CH₂CF₃, or OCH₂CH₂CH₂CF₃; R₂ = perfluoroalkyl; R₃ = H or halo; X = OH, alkoxy, alkylthio, imino, alkylimino, halo, etc.; Z = N or CH] or salts, N-oxides, or tautomers thereof are prepared as neoplastic inhibitors for the treatment of human or animal body. For example, the compound II was prepared in a multi-step synthesis. Formulations containing I as an active ingredient were also described.

IT 709045-02-3P
 RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
 (drug candidate, reactant; preparation of anthranilic acid amide derivs. as neoplastic inhibitors)

RN 709045-02-3 HCAPLUS

CN Benzamide, 2-[[[6-methoxy-5-phenyl-3-pyridinyl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



IT 709045-03-4P 709045-42-1P 709045-43-2P
 709045-44-3P 709045-45-4P 709045-48-7P
 709045-49-8P 709045-50-1P 709045-51-2P
 709045-52-3P 709045-53-4P 709045-54-5P
 709045-55-6P 709045-56-7P 709045-57-8P
 709045-58-9P 709045-59-0P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

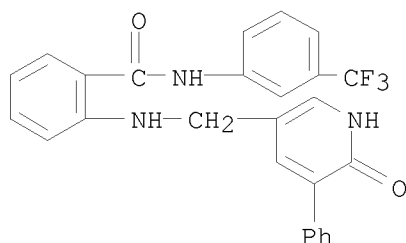
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(drug candidate; preparation of anthranilic acid amide derivs. as neoplastic inhibitors)

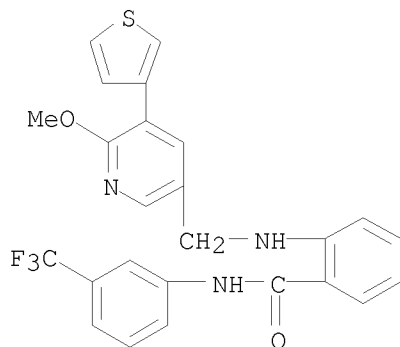
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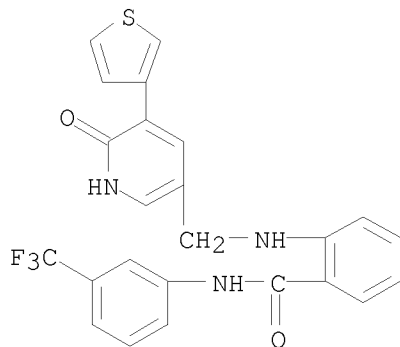
RN 709045-42-1 HCAPLUS

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RN 709045-43-2 HCAPLUS

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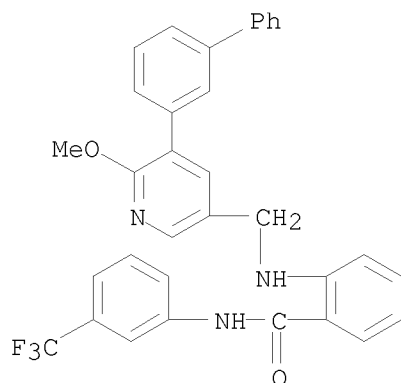


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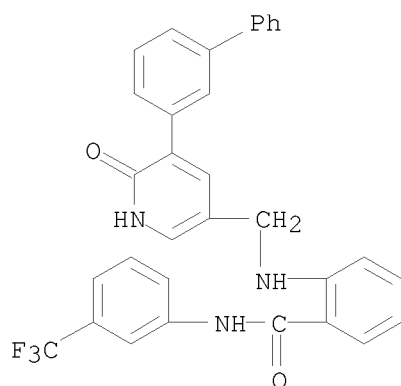
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RN 709045-45-4 HCAPLUS

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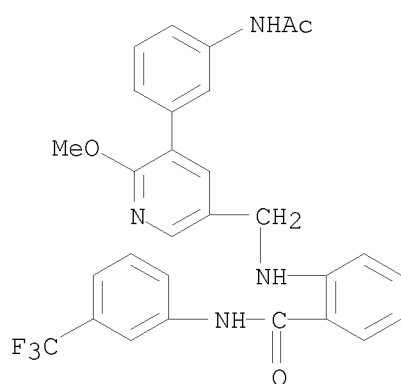


RN 709045-48-7 HCAPLUS

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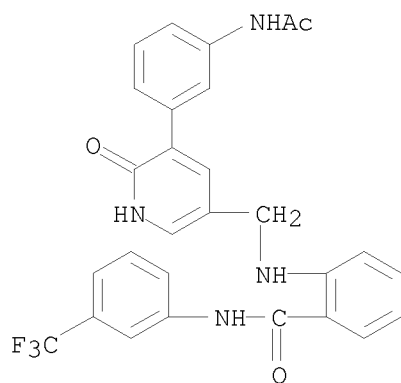
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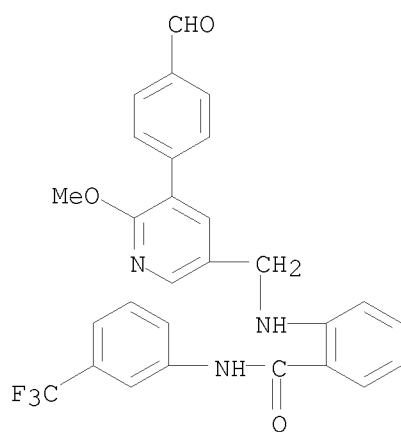
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RN 709045-50-1 HCAPLUS

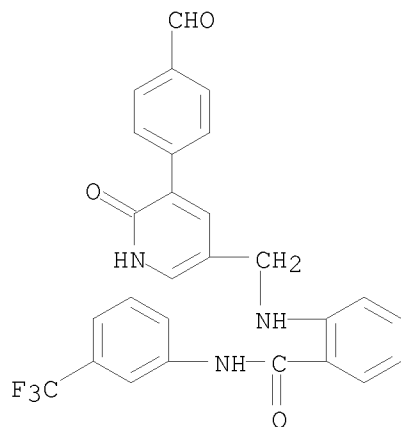
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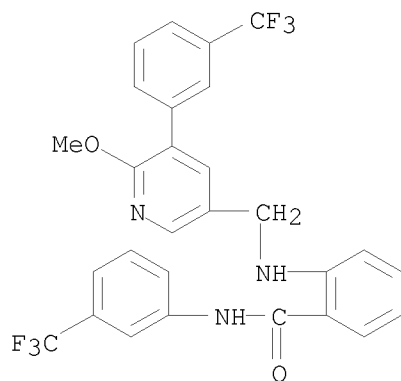
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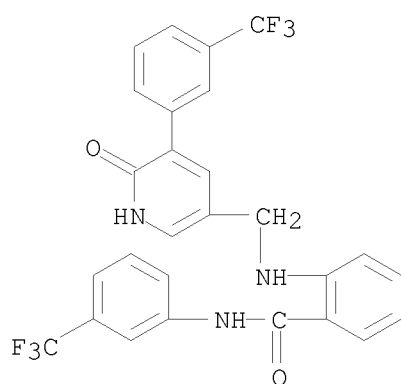
RN 709045-52-3 HCAPLUS
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RN 709045-53-4 HCAPLUS
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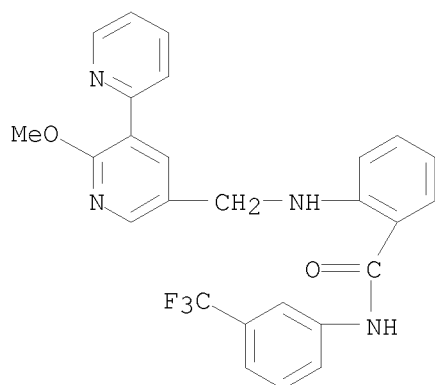
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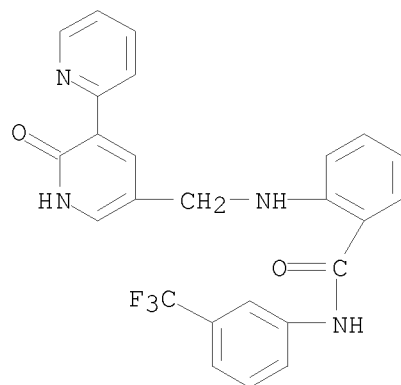
RN 709045-54-5 HCAPLUS

CN Benzamide, 2-[[[(2'-methoxy[2,3'-bipyridin]-5'-yl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



RN 709045-55-6 HCAPLUS

CN Benzamide, 2-[[[(1',2'-dihydro-2'-oxo[2,3'-bipyridin]-5'-yl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)

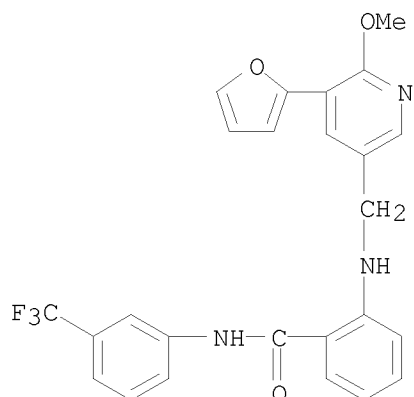


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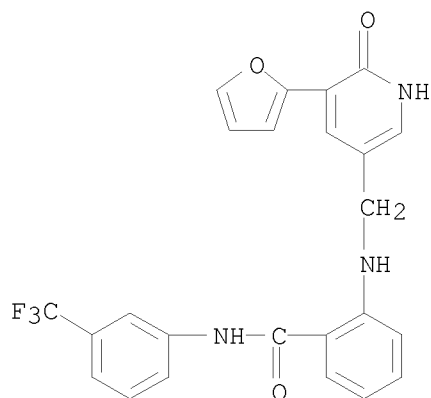
RN 709045-56-7 HCAPLUS

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RN 709045-57-8 HCAPLUS

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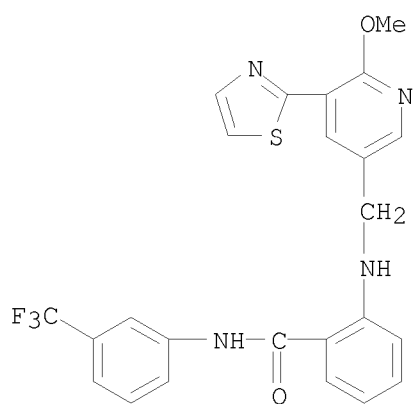


RN 709045-58-9 HCAPLUS

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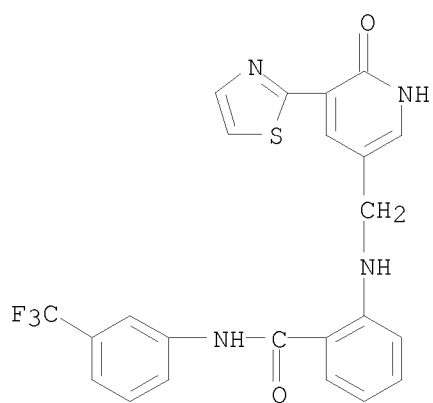
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STN Search



RN 709045-59-0 HCAPLUS

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